

<https://helda.helsinki.fi>

---

# NON-INVASIVE VASCULAR VERY-HIGH RESOLUTION ULTRASOUND TO QUANTIFY ARTERY INTIMA LAYER THICKNESS : VALIDATION OF THE FOUR-LINE PATTERN

Sundholm, Johnny K. M.

2019-08

---

Sundholm , J K M , Paetau , A , Albäck , A , Pettersson , T & Sarkola , T 2019 , '  
NON-INVASIVE VASCULAR VERY-HIGH RESOLUTION ULTRASOUND TO QUANTIFY  
ARTERY INTIMA LAYER THICKNESS : VALIDATION OF THE FOUR-LINE PATTERN ' ,  
Ultrasound in Medicine & Biology , vol. 45 , no. 8 , pp. 2010-2018 . <https://doi.org/10.1016/j.ultrasmedbio.2019.04.017>

---

<http://hdl.handle.net/10138/316310>

<https://doi.org/10.1016/j.ultrasmedbio.2019.04.017>

---

cc\_by\_nc\_nd

acceptedVersion

---

*Downloaded from Helda, University of Helsinki institutional repository.*

*This is an electronic reprint of the original article.*

*This reprint may differ from the original in pagination and typographic detail.*

*Please cite the original version.*

Sundholm Assessing Vascular Intima with Ultrasound

## Non-invasive vascular very-high resolution ultrasound to quantify artery intima layer thickness: validation of the four-line pattern

Johnny K.M. Sundholm, MD<sup>1</sup>

Anders Paetau, MD, PhD<sup>2</sup>

Anders Albäck, MD, PhD<sup>3</sup>

Tom Pettersson, MD, PhD<sup>4</sup>

Taisto Sarkola, MD, PhD<sup>1</sup>

<sup>1</sup>Children's Hospital, University of Helsinki and Helsinki University Hospital, Helsinki, Finland

<sup>2</sup> HUSLAB Meipa1 pathology laboratory & Haartman Institute, University of Helsinki, Helsinki, Finland

<sup>3</sup> Department of Vascular Surgery, University of Helsinki and Helsinki University Hospital, Helsinki, Finland

<sup>4</sup> Department of General Internal Medicine and Geriatrics, University of Helsinki and Helsinki University Hospital, Helsinki, Finland

Abstract word count 169, Figures 3, Tables 2, References 32, Supplemental tables 6, Supplemental figures 4.

Address for Correspondence: Johnny Sundholm, MD, Children's Hospital, Helsinki University Hospital, Stenbäckinkatu 11, POB 281, FIN-00029, Helsinki, Finland, Phone: +358504011154, Fax: +358947174706, e-mail: johnny.sundholm@helsinki.fi

**Abstract**

Preliminary findings suggest that very-high resolution ultrasound (VHRU, 55 MHz) could differentiate arterial intima layer thickness (IT) non-invasively *in vivo*. We aimed to validate ultrasound derived IT measurements and describe a four-line pattern consistent with intimal thickening. VHRU was applied to temporal arteries of 37 patients with suspected giant cell arteritis without inflammation on histology. Anatomically matched ultrasound derived measurements of arterial layer thickness with the leading-edge method was compared to histology. Intimal thickening (IT>0.06 mm on histology) was identified as a four-line pattern in VHRU with a sensitivity of 96.3% and a specificity of 100%. Histological and VHRU IT measurement agreement was excellent (mean difference 0.007mm; 95%LOA - 0.043-0.057) and ICC 0.923 (95%CI 0.833-0.964). Intra- and inter-observer agreements for VHRU IT was high: ICC 0.946 (95%CI 0.877-0.976) and 0.872 (95%CI 0.773-0.943). VHRU utilizing the leading-to-leading edge method allows accurate and reliable measurements of arterial IT in patients with IT>0.06mm. Measurements of IT will provide the opportunity to explore early subclinical structural intimal changes in the arterial wall increasing with age.

Keywords: Vascular imaging, intima thickness, intima-media thickness, ultrasound biomicroscopy, methodology, very-high resolution ultrasound

## 1 Introduction

2 Intimal thickening is a key process in vascular aging related to atherosclerosis and  
3 increased cardiovascular mortality.(Enos et al. 1953; McNamara et al. 1971; Mönckeberg  
4 1915; O'Leary et al. 1999) In 1984 Pignoli et al were among the first to introduce non-  
5 invasive measurements of vascular wall-thickness using B-mode high-resolution ultrasound  
6 (HRU).(Pignoli et al. 1984, 1986) In their landmark paper they compared ultrasound images  
7 of the carotid artery with histology and described the double line pattern of the arterial far  
8 wall. Since then, carotid intima-media thickness (IMT) has been shown to be an independent  
9 predictor of cardiovascular risk and used as a surrogate marker for cardiovascular  
10 disease.(Burke et al. 1995; Lorenz et al. 2018; Stein et al. 2008) In the aging population, an  
11 increased IMT has been attributed mainly to intimal thickening with differences between  
12 arteries and even between locations in the same artery.(Chowdhury et al. 2004; Nakashima  
13 et al. 2002; Wilens 1951)

14 Due to the physical properties of ultrasound the echogenic region is created at the  
15 interface of two different mediums with the leading edge defining the true tissue border  
16 whereas the trailing edge is unrelated to the measured dimension, mainly related to  
17 ultrasound frequency.(Sarkola et al. 2010; Wendelhag et al. 1991) As a result, IMT needs to  
18 be measured leading edge to leading edge from the vascular far wall for the measurements  
19 to be accurate.(Wikstrand 2007) The measurement of arterial intima layer thickness (IT),  
20 arterial wall layer thicknesses in small peripheral arteries, and arteries in the pediatric  
21 population is, thus, limited using conventional HRU frequencies due to insufficient ultrasound  
22 resolution.(Foster et al. 1993; Sarkola et al. 2010; Sundholm et al 2015)

23 Very-high resolution ultrasound (VHRU, 25-55MHz, axial resolution 0.105-0.045mm)  
24 provides the opportunity to image vascular structures in more detail *in vivo*, limited mainly by  
25 penetration depth. VHRU is able to assess the IMT of peripheral conduit muscular arteries,  
26 and the adventitial thickness (AT), seen as a triple line pattern, with additional echolucent  
27 and echogenic zones.(Sarkola et al. 2010) The non-invasive measurement of IT with very-

high resolution ultrasound frequencies, similar to the use of intravascular ultrasound,(Siegel et al. 1993) has previously been validated using a leading-to-trailing edge method applied to the first echogenic zone of the far wall blood-intima interface.(Choi et al. 2009; Dangardt et al. 2018; Osika et al. 2007; Rodriguez-Macias et al. 2001) However, the leading-to-trailing edge method of IT has lately been questioned in validation studies using healthy arterial specimens. It has been shown that VHRU is unable to quantify intimal thickness in subjects with an intimal thickness below ultrasound axial resolution (as reviewed in Sarkola et al. 2010 discussion). In a more recent study a distinct four-line pattern with an additional echolucent zone separating the first echogenic zone of the blood-intima interface into three separate zones has been described in relation to radiotherapy among long-term childhood cancer survivors.(Vatanen et al. 2015) The study interpreted the finding, a four-line pattern, as a thickened intima and a sign of early vascular aging but without histological verification of the measurement.

The aim of this study was to validate VHRU derived IT measurements in comparison to histology, and to explore the distinct four-line pattern observed in the VHRU image. Our hypothesis was that the additional region in the distinct four-line pattern corresponds to the IT in arteries with intimal thickening. We further explored the presence of the four-line pattern in relation to age and cardiovascular risk factors.

## **Material and Methods**

We prospectively recruited 74 consecutive patients with suspected giant cell arteritis (GCA) referred to the unit of Vascular Surgery for temporal artery biopsy as part of routine diagnostics between 8/2015 and 2/2018. Exclusion criteria were any sign of inflammation on histology, missing histology or diagonally sliced histology precluding histological measurements, and subjects without 55MHz VHRU image. Subject characteristics were recorded at presentation using a standard questionnaire filled-in by the investigator and by reviewing patient records to assess background information on cardiovascular risk including smoking history, lipid disorders, diabetes, and hypertension, and current medications. The

local ethics board and the hospital approved the research protocol, and written consent was obtained from the participants.

### *Ultrasound systems*

Two different ultrasound systems were used in this study, the Vevo 770 (VisualSonics, Toronto, ON, Canada) for the first 41 patients and, due to malfunction of the Vevo 770, Vevo MD (VisualSonics, Toronto, ON, Canada) for the following 33 patients. The Vevo 770 was equipped with mechanical RMV-710B, RMV-712, RMV-708 transducers with center frequencies 25, 35 and 55MHz respectively. The Vevo MD was equipped with electronic UHF22, UHF48 and UHF70 transducers, with 15, 30 and 50MHz center frequencies respectively. Further comparison of the ultrasound systems is available in Supplemental table 1.

### *VHRU IT validation*

Vascular ultrasound images were obtained 1h prior to biopsy by one skilled investigator (JS) using a 55 MHz transducer (axial resolution 15.4 $\mu$ m/pixel) with the Vevo 770 system, and a UHF70 transducer (axial resolution 12.3 $\mu$ m/pixel) with Vevo MD. Superficial temporal arteries were screened bilaterally for pathology proximal to the vessel bifurcation using VHRU. Scanning area was marked on the skin at the end of imaging to guide the surgeon and to ensure the biopsy was matched to the site of imaging. IT and IMT was measured by one skilled investigator (JS) from the images offline with manufacturer supplied vendor software Vevo 3.0.0 (Vevo 770) and VevoLab 2.0.0 (Vevo MD) in end-diastole using both far wall leading-to-leading edge and leading-trailing edge methods with electronic calipers prior to histological processing. Measurements were done at the central section of the VHRU images as the biopsy would be sectioned from the middle of the specimen. The mean of three measurements was used in analyses. Measurements for a subset (n=31) of images were independently repeated after 1-month by the first investigator to assess intra-observer agreement. Inter-observer agreement was assessed by

independent measurements performed on the same subset by a second skilled investigator (TS). Image quality was subjectively evaluated in all images and grouped into three quality classes: 3. Highest quality, with the far wall visible across the entire images, 2. moderate quality, the far wall visible across most of the images, 1. low quality, with the far wall visible in only a small part of the image.

### *Biopsy and histology*

The biopsy procedure was performed as routine GCA diagnostics at the Helsinki University Central Hospital Department of Vascular Surgery. The biopsy specimens were fixed in formalin and cut in transverse sections, stained with hematoxylin and eosin (H&E) stain and Verhoff's elastic stain (VEG).(Alturkistani et al. 2015) Biopsies were evaluated for vascular pathology at a certified pathology unit (HUSLAB). Only negative biopsies without any signs of vascular inflammation were included in this study.(Cavazza et al. 2014) Twenty specimens were positive for GCA, three biopsies did not include arterial structures and eight of the histological specimens were sectioned diagonally precluding comparable histological measurements with VHRU arterial layer thickness. VHRU images of sufficient resolution were missing in six subjects. Remaining 37 subjects were included in the study. Biopsy specimens were sectioned in transverse section. Histological sections were evaluated using optic microscopy, photographed at 10x zoom (Nikon Eclipse 80i & Digital Sight DS-5M, Tokyo, Japan) and IT and IMT later measured offline using ImageJ 1.51J8 (National Institutes of Health, Bethesda, MD, USA)(Schneider et al. 2012) with electronic calipers, as a mean of 10 different measurements, covering the whole circular vascular wall to avoid error due to local differences in vascular wall dimensions.

### *VHRU imaging of peripheral conduit arteries*

The presence of four-line pattern was also assessed in peripheral conduit arteries in a majority of the GCA cohort with VHRU. Peripheral conduit artery imaging was not performed in some subjects due to time constraint at the research appointment. Images

were obtained from bilateral common carotid arteries, the right radial, and right brachial artery. Carotid images (n=30) were obtained, according to guidelines(Stein et al. 2008), 1cm proximal to the carotid bulb using the 25 MHz (Vevo 770) or UHF22 (Vevo MD) transducers. Common carotid artery images were evaluated for plaque presence throughout the carotid bulb and the proximal parts of the external carotid artery. Plaques were defined according to the Mannheim consensus as focal thickening of IMT fulfilling one of the following criteria: 1.  $IMT > 1.5mm$ , or 2. IMT increase of  $0.5mm > 50\%$  compared to the surrounding IMT.(Touboul et al. 2012) Plaque burden was further graded according to plaque number as single or multiple plaques. Radial artery images (n=34) were acquired 1cm proximal to the *palma manus* using the 55 MHz (Vevo 770) or UHF70 (Vevo MD) transducers, and brachial arteries (n=28) were imaged 3cm proximal to the cubital skin fold using 35 MHz (Vevo 770) or UHF48 (Vevo MD) transducers. IMT and intima-media-adventitia thickness (IMAT) were measured from the far wall using the leading-edge method in end-diastole.(Sarkola et al. 2010)

#### *Radial artery four-line pattern in relation to age*

To broaden the age spectrum and investigate presence of four-line pattern over age, we combined our GCA sample (age 40-86, n=64) with a convenience sample of subjects recruited as part of other projects (final sample n=444). The convenience sample consisted of 1. healthy children and adolescents (age 0-18 yrs, n=139, previously published study (Sarkola et al. 2012)), 2. teenagers with type 1 diabetes (age 13-16 yrs, n=39, unpublished), 3. healthy males (age 20-46 yrs, n=24, unpublished), and 4. women with obesity and or gestational diabetes (age 28-51 yrs, n=178, unpublished). The convenience sample is further described in supplemental table 2. To assess IMT and the presence of the four-line pattern (IT) in relation to age, VHRU images of the radial artery were obtained with 55 MHz (or UHF70) and assessed as mentioned above.

#### *Data analysis*



Results are reported as mean with SD, median with minimum and maximum values, and ratios, as appropriate. Student's T-test was used for comparison between groups. Comparisons between independently obtained ultrasound and histology measurements as well as intra- and inter-observer agreements were quantified by calculating the mean difference, 95% limits of agreement, coefficient of variation (CV) and intraclass correlation coefficient (ICC). Ultrasound and histology measurements were further compared using Bland-Altman plots.(Bland and Altman 1986) The diagnostic utility of intimal thickening using the presence of four-line pattern was evaluated using receiver operating characteristics (ROC) analysis. Multiple linear regression was used to assess the effects of age and cardiovascular risk factors on histological IT. The occurrence of four-line pattern in relation to age and radial artery intima-media thickness was assessed using logistic regression. Statistical analysis was done using SPSS version 24 (IBM, Armonk, NY, USA) and STATA MP 15.1 (StataCorp LCC, College Station, TX, USA).

## Results

37 patients were included in the validation study (Vevo 770 n=20, and Vevo MD n=17). Patient mean age was 67 (SD 9,8) years, there were 24 (65%) females, 4 (11%) subjects reported current daily and 16 (43%) previous daily smoking, 17 (46%) were treated for hypertension, 9 (24%) for type 2 diabetes, 13 (35%) for hypercholesterolemia, and 4 (11%) had a history of coronary artery disease in angiography.

### *Validation of VHRU IT and the four-line pattern*

In temporal artery VHRU images intimal thickening was observed as a separation of the traditional first echogenic zone into three separate zones (1-3). Compared to histology, the four-line pattern of the far wall in VHRU images consisted of the blood-intima interface (zone 1), echolucent intima core (zone 2), echodense IEL (internal elastic lamina), i.e. intima-media interface (zone 3), echolucent media layer (zone 4), echodense EEL (external

elastic lamina), i.e. media-adventitia interface (zone 5), echolucent adventitia (zone 6) and echodense adventitia-extravascular fat interface (zone 7) (Figure 1a and b) . In the setting of a thin intima the first 3 echo zones (zones 1-3 in figure 1c) were fused. The adventitial thickness of the temporal artery was bordering the resolution of the VHRU method (mean thickness 0.06mm +/- 0.02mm) leading to a fusion of the echo zones 5-7 in most images, as seen in figure 1c. The far wall four-line pattern in VHRU images of the temporal artery (Figure 1b) was similar to the far wall VHRU images of an aged muscular radial artery (Figure 2c), consistent with a thickened intima layer in the radial artery.

The mean histological temporal artery IT in the sample was 0.10mm (SD: 0.05; range: 0.02-0.22mm, Supplemental figure 1). Temporal artery four-line pattern, i.e. measurable IT, was identifiable in 28 patients (76%) using VHRU. The mean histological IT in images with no four-line pattern on VHRU was 0.04mm (SD: 0.04; range 0.02-0.12mm), with 8/9 arteries having an intimal thickness less than 0.06mm. ROC-analysis of four-line presence and IT on histology gave an AUC of 0.99 (95%CI: 0.97-1.00) with the 4-line pattern predicting a histological IT>0.06mm with a sensitivity of 96.3% (95%CI: 81.0-99.9%) and a specificity of 100.0% (95%CI: 66.3-100.0%). All images were of high or moderate quality (quality classes 3 and 2, n=17 and n=20, respectively).

Ultrasound measurements of IT and IMT agreed with histological measurements (Table 1 and Figure 3) confirming method accuracy. Intra- and inter-observer agreement of the IT and IMT measurements were good (Table 2 and Figure 3). We further compared the histological agreements using the different ultrasound devices Vevo 770 and the Vevo MD (Supplemental table 3), both agreeing with histology. The Vevo MD had slightly higher ICC compared to the Vevo 770 (ICC 0.867, 95%CI 0.603-0.955 and ICC 0.971 95%CI 0.901-0.992) for IT measurements, a difference mainly related to an outlier in the Vevo 770 sample (ICC 0.931 95%CI 0.785-0.978 after exclusion of the outlier), with no change in accuracy over time suggesting that the differences were unrelated to a learning curve. Images of

higher quality (quality class 3) had slightly higher accuracy than those with moderate quality (quality class 2) with CV% 12.8 and 23.6  $p=0.098$ , respectively.

The leading-to-trailing edge VHRU measurements of blood-intima interface (fused zone 1-3) in arteries with thin ( $<0.06\text{mm}$ ) IT consistently overestimated histological IT (mean difference  $0.018\text{mm}$  95%CI LOA  $-0.004-0.032\text{mm}$ ) with bias related to the histological IT level (Supplemental figure 2a). Furthermore, leading-to-trailing edge VHRU measurements of IT (zones 1-3) in arteries with visible intimal thickening systematically overestimated IT (mean difference  $0.051$ , 95%LOA  $0.019-0.084$ ), with constant bias in the measurement over the range of histological IT (Supplemental figure 2b). The bias in the leading-to-trailing edge VHRU measurement is further clarified in Supplemental figure 3.

In a linear regression model temporal artery histological IT [ $\mu\text{m}$ ] was significantly predicted by age [years] ( $\beta=1.7$   $p=0.038$ ), hypertension [y/n] ( $\beta=42.2$   $p=0.020$ ), diabetes [y/n] ( $\beta=34.2$   $p=0.044$ ), but not by smoking [10 pack years] ( $\beta=6.8$   $p=0.069$ ), or hypercholesterolemia [y/n] ( $\beta=-26.7$   $p=0.141$ ). The adjusted  $R^2$  of the model was  $0.404$ ,  $p<0.001$  (supplemental table 4).

#### *In vivo peripheral arterial layer thickness and presence of plaques in GCA group.*

Carotid screening was feasible using the 25MHz in all 30 subjects, whereas the 35MHz transducer had insufficient penetrance to clearly view the far wall of the carotid artery. 40% of the subjects had identifiable plaques in the carotid arteries, with a majority having multiple plaques. None of the carotid arteries showed separation of the first echogenic zone. The 55MHz transducer had sufficient penetrance to image the radial artery in all subject and there was an identifiable four-line pattern in 68% of the subjects. The deeper situated brachial artery required the 35MHz transducer (and 25MHz in 3 subjects) to view the vascular far wall. The four-line pattern was identifiable in only one (4%) brachial artery (supplemental table 5.)

#### *Intimal thickening observed with VHRU in relation to age*

In a convenience sample of radial artery VHRU images IMT increased with age and the four-line pattern was identified in 51/444 (11.7%) subjects aged >36yrs (supplemental figure 4.), but not in younger subjects. The presence of the four-line pattern was independently predicted by both age ([10yrs] OR 3.077; CI95% 2.125-4.456;  $p<0.001$ ) and IMT ([0.1mm] OR 4.629; CI95% 1.631-13.140;  $p=0.004$ ) in a logistic regression model (supplemental table 6). The feasibility of radial artery VHRU was excellent and adequate imaging was obtained in all patients attempted.

## Discussion

The aim of this study was to validate the vascular non-invasive VHRU method with histology to assess whether the leading-to-leading-edge measurement of the first echolucent region in the four-line pattern corresponds to the IT of the artery. The data shows that the VHRU method provides an accurate and reliable measurement of IT in superficial arteries with layer thickness exceeding axial ultrasound resolution, and that the presence of a four-line pattern is consistent with  $IT>0.06\text{mm}$ . This method provides the opportunity to investigate the vascular health in the aging population noninvasively *in vivo*.

We further show that intimal thickness increases with age and cardiovascular risk factors, and that the four-line pattern is present in peripheral muscular arteries in a substantial proportion (76%) of aging subjects, even more prevalent than carotid plaques. Whereas there was a visible four-line pattern in a majority of the temporal and radial arteries in the GCA cohort, it was absent in all but one brachial artery. This might be due to the lower frequency used or differences in the vascular aging process between the arteries.

There have been previous attempts to validate the IT measurement using non-invasive VHRU. Rodriguez-Macias et al (2001) reported systematic overestimation of ultrasound derived IT using 25 MHz. Moreover, Osika et al (2007) compared ultrasound images of the human mesenteric artery obtained with 55 MHz with histology (IT ranging from 100-400 $\mu\text{m}$ ) and showed a good correlation between measurements. The studies

implemented the method on *in vivo* images measuring the thickness of the echogenic zone of the blood-intima interface with the leading-to-trailing edge method in muscular and carotid arteries suggesting that it represents the intimal layer.(Osika et al. 2007; Rodriguez-Macias et al. 2001) Sarkola et al (2010) found no correlation between the thickness of the echogenic blood-intima interface and histological intima thickness in histology in healthy arterial specimens with thin IT obtained from rabbits and pigs and further reported that the measured interface thickness increased with decreasing ultrasound frequency, and concluded that histological IT was beyond VHRU resolution in healthy arteries.

The IT of healthy arteries, consisting of a layer of endothelial cells and the internal elastic lamina, typically measures 10-30 $\mu$ m as shown by histological data.(Kölliker 1854; Velican and Velican 1981) The healthy intima is beyond ultrasound axial resolution even for VHRU. The theoretical minimum measurable thickness of a vascular layer using the leading-edge method is three pixels, i.e.  $3 \times 0.0123 = 0.0369$ mm for the UHF70 transducer and slightly higher for the 55MHz transducer. Non-invasive imaging at this level requires optimal conditions, and in our experience, the layer thickness quantifiable with an acceptable level of certainty is somewhat higher (~5px, i.e. 0.06 mm). The measurement of arterial IT is dependent on ultrasound frequency, vessel size and location due to reduced penetrance of the higher frequencies used. We did not find any consistent difference in accuracy between the two ultrasound systems, with no considerable difference in axial resolution. Further, age related changes in the vascular wall including fragmentation of the IEL and increasing heterogeneity of the wall structure is inevitably related to reduced image quality, seen as a trend for reduced accuracy not reaching significance in our sample.

Our results confirm previous validations for ultrasound derived IMT and shows that the far wall four-line pattern of the VHRU image indicates an intimal layer with a quantifiable thickness. Furthermore, we show that the leading-to-leading edge method provides a more accurate measure of intimal thickness than the leading-to-trailing edge method applied in previous IT validations. These findings using human arterial specimens confirm a systematic

overestimation of IT using the leading-to-trailing edge method to assess the blood-intima interphase in healthy vessels, with a very thin intima on histology, as previously shown.(Sarkola et al. 2010) Our findings demonstrate that in arteries with thin IT, beyond the axial resolution of the method, the IT consistently correspond to approximately three pixels, and thus giving a bias equal to three pixels minus histological IT (mean difference 0.018 mm), whereas in arteries, with a visible separation of the intima from the intima-media complex, incorporation of the trailing edge includes part of the media, again, consistent with three pixels (mean difference 0.051mm, supplemental figure 2). The present results support recommendations that leading-to-leading edge measurements should be used to quantify intimal thickening displayed as a four-line pattern in the VHRU image similar to the far wall arterial IMT measurement.(Wendelhag et al. 1991)

The main limitation of this study is that arterial specimens from temporal arteries were included only. However, histologically the temporal artery is a muscular artery with a morphology resembling that of other peripheral conduit arteries that may be imaged with VHRU frequencies. The results should therefore be applicable for the assessment of arterial IT. Another limitation is that the subjects were recruited from patients remitted for TA biopsy due to suspected GCA. We did exclude patients with any sign of vascular inflammation, and vascular changes should mainly be related to cardiovascular risk factors. The prevalence of cardiovascular risk factors in our subjects was similar to the average elderly population, suggesting applicability of this method in this age group.(Jain and Paranjape 2013; Lionakis et al. 2012) A further limitation is that we used two different ultrasound systems without cross validation. The transducers used are almost similar in regards of resolution and penetrance, and both agreed well with histology. The strength of our study is the histological comparison to anatomically matched *in vivo* imaging of a relatively large sample of arterial specimens with a range of intimal thickening.

*Conclusion*

1           We present a novel non-invasive very-high resolution ultrasound-based method for  
2 the quantification of thick IT of superficial arteries in the aging population. The measurement  
3 is based on identification of a four-line pattern of the arterial far wall appearing in arteries  
4 with a thickened intima layer ( $>0.06\text{mm}$ ). Our results support the use of the leading-to-  
5 leading edge method of the arterial far wall in layer thickness quantification. The leading-to-  
6 trailing edge method systematically overestimated histological layer thickness. This  
7 technique will provide the opportunity to explore early subclinical atherosclerotic changes in  
8 the arterial wall increasing with age.

9 *Sources of funding*

10           This study has been supported by grants from the Sigrid Juselius Foundation, The  
11 Medical Society of Finland, and Finnish Foundation for Pediatric Research, Perklen  
12 foundation, Medicinska understödsföreningen Liv och Hälsa, and the Stockmann  
13 Foundation.

14 *Conflict of interest*

15           The authors declare that there is no conflict of interest.

## 1   References

- 2   Alturkistani HA, Tashkandi FM, Mohammedsaleh ZM. Histological Stains: A Literature  
3   Review and Case Study. Glob J Health Sci 2015;8:72-9.
- 4   Bland JM, Altman DG. Statistical methods for assessing agreement between two methods of  
5   clinical measurement. Lancet 1986;1:307-10.
- 6   Burke GL, Evans GW, Riley WA, Sharrett AR, Howard G, Barnes RW, Rosamond W, Crow  
7   RS, Rautaharju PM, Heiss G. Arterial wall thickness is associated with prevalent  
8   cardiovascular disease in middle-aged adults. The Atherosclerosis Risk in Communities  
9   (ARIC) Study. Stroke 1995;26:386-91.
- 10   Cavazza A, Muratore F, Boiardi L, Restuccia G, Pipitone N, Pazzola G, Tagliavini E, Ragazzi  
11   M, Rossi G, Salvarani C. Inflamed temporal artery: histologic findings in 354 biopsies, with  
12   clinical correlations. Am J Surg Pathol 2014;38:1360-70.
- 13   Choi Y, Youn H, Youn J, Park C, Oh Y, Chung W. Measurement of the Intimal Thickness of  
14   the Carotid Artery: Comparison Between 40 MHz Ultrasound and Histology in Rats.  
15   Ultrasound Med Biol 2009;35:962-6.
- 16   Chowdhury UK, Airan B, Mishra PK, Kothari SS, Subramaniam GK, Ray R, Singh R,  
17   Venugopal P. Histopathology and morphometry of radial artery conduits: basic study and  
18   clinical application. Ann Thorac Surg 2004;78:1614-21.
- 19   Dangardt F, Charakida M, Chiesa S, Bhowruth D, Rapala A, Thurn D, Schaefer F, Deanfield  
20   J, Shroff R. Intimal and medial arterial changes defined by ultra-high-frequency ultrasound:  
21   Response to changing risk factors in children with chronic kidney disease. PLoS One  
22   2018;13:e0198547.



- 1 Enos WF, Holmes RH, Beyer J. Coronary disease among United States soldiers killed in  
2 action in Korea; preliminary report. J Am Med Assoc 1953;152:1090-3.
- 3 Foster FS, Lockwood GR, Ryan LK, Harasiewicz KA, Berube L, Rauth AM. Principles and  
4 applications of ultrasound backscatter microscopy. IEEE Trans Ultrason Ferroelectr Freq  
5 Control 1993;40:608-17.
- 6 Jain A, Paranjape S. Prevalence of type 2 diabetes mellitus in elderly in a primary care  
7 facility: An ideal facility. Indian J Endocrinol Metab 2013;17:S318-22.
- 8 Kölliker A. Of The Blood Vessels: Arteries. In: Busk G, Huxley T, eds. **Manual of Human**  
9 **Histology**. 2nd edition. London: The Sydenham Society, 1854:294,295-301.
- 10 Lionakis N, Mendrinou D, Sanidas E, Favatas G, Georgopoulou M. Hypertension in the  
11 elderly. World J Cardiol 2012;4:135-47.
- 12 Lorenz MW, Gao L, Ziegelbauer K, Norata GD, Empana JP, Schmidtman I, Lin HJ,  
13 McLachlan S, Bokemark L, Ronkainen K, Amato M, Schminke U, Srinivasan SR, Lind L,  
14 Okazaki S, Stehouwer CDA, Willeit P, Polak JF, Steinmetz H, Sander D, Poppert H,  
15 Desvarieux M, Ikram MA, Johnsen SH, Staub D, Sirtori CR, Iglseder B, Beloqui O, Engstrom  
16 G, Frier A, Rozza F, Xie W, Parraga G, Grigore L, Plichart M, Blankenberg S, Su TC,  
17 Schmidt C, Tuomainen TP, Veglia F, Volzke H, Nijpels G, Willeit J, Sacco RL, Franco OH,  
18 Uthoff H, Hedblad B, Suarez C, Izzo R, Zhao D, Wannarong T, Catapano A, Ducimetiere P,  
19 Espinola-Klein C, Chien KL, Price JF, Bergstrom G, Kauhanen J, Tremoli E, Dorr M,  
20 Berenson G, Kitagawa K, Dekker JM, Kiechl S, Sitzer M, Bickel H, Rundek T, Hofman A,  
21 Mathiesen EB, Castelnovo S, Landeche MF, Rosvall M, Gabriel R, de Luca N, Liu J,  
22 Baldassarre D, Kavousi M, de Groot E, Bots ML, Yanez DN, Thompson SG, PROG-IMT  
23 study group. Predictive value for cardiovascular events of common carotid intima media  
24 thickness and its rate of change in individuals at high cardiovascular risk - Results from the  
25 PROG-IMT collaboration. PLoS One 2018;13:e0191172.

- 1 McNamara JJ, Molot MA, Stremple JF, Cutting RT. Coronary artery disease in combat  
2 casualties in Vietnam. JAMA 1971;216:1185-7.
- 3 Mönckeberg J. Über die Atherosklerose der Kombattanten (nach Obduktionsbefunden).  
4 Zentralbl Herz Gefäßkrankheiten 1915:7-10.
- 5 Nakashima Y, Chen Y, Kinukawa N, Sueishi K. Distributions of diffuse intimal thickening in  
6 human arteries: preferential expression in atherosclerosis-prone arteries from an early age.  
7 Virchows Archiv 2002;441:279-88.
- 8 O'Leary DH, Polak JF, Kronmal RA, Manolio TA, Burke GL, Wolfson SK, Jr. Carotid-artery  
9 intima and media thickness as a risk factor for myocardial infarction and stroke in older  
10 adults. Cardiovascular Health Study Collaborative Research Group. N Engl J Med  
11 1999;340:14-22.
- 12 Osika W, Dangardt F, Grönros J, Lundstam U, Myredal A, Johansson M, Volkmann R,  
13 Gustavsson T, Ming Gan L, Friberg P. Increasing Peripheral Artery Intima Thickness From  
14 Childhood to Seniority. Arterioscler Thromb Vasc Bio 2007;27:671-6.
- 15 Pignoli P. Ultrasound B-mode imaging for arterial wall thickness measurement.  
16 Atherosclerosis Reviews 1984:177-84.
- 17 Pignoli P, Tremoli E, Poli A, Oreste P, Paoletti R. Intimal plus medial thickness of the arterial  
18 wall: a direct measurement with ultrasound imaging. Circulation 1986;74:1399-406.
- 19 Rodriguez-Macias KA, Naessen T, Bergqvist D. Validation of in vivo noninvasive high-  
20 frequency ultrasonography of the arterial wall layers. Ultrasound Med Biol 2001;27:751-6.
- 21 Sarkola T, Manhiot C, Slorach C, Bradley TJ, Hui W, Mertens L, Redington A, Jaeggi E.  
22 Evolution of the arterial structure and function from infancy to adolescence is related to

anthropometric and blood pressure changes. *Arterioscler Thromb Vasc Biol* 2012;32:2516-24.

Sarkola T, Redington A, Keeley F, Bradley T, Jaeggi E. Transcutaneous very-high-resolution ultrasound to quantify arterial wall layers of muscular and elastic arteries: Validation of a method. *Atherosclerosis* 2010;212:516-23.

Schneider CA, Rasband WS, Eliceiri KW. NIH Image to ImageJ: 25 years of image analysis. *Nat Methods* 2012;9:671-5.

Siegel RJ, Chae JS, Maurer G, Berlin M, Fishbein MC. Histopathologic correlation of the three-layered intravascular ultrasound appearance of normal adult human muscular arteries. *Am Heart J* 1993;126:872-8.

Stein JH, Korcarz CE, Hurst RT, Lonn E, Kendall CB, Mohler ER, Najjar SS, Rembold CM, Post WS, American Society of Echocardiography Carotid Intima-Media Thickness Task Force. Use of carotid ultrasound to identify subclinical vascular disease and evaluate cardiovascular disease risk: a consensus statement from the American Society of Echocardiography Carotid Intima-Media Thickness Task Force. Endorsed by the Society for Vascular Medicine. *J Am Soc Echocardiogr* 2008;21:93,111; quiz 189-90.

Sundholm JKM, Olander RFW, Ojala TH, Andersson S, Sarkola T. Feasibility and precision of transcutaneous very-high resolution ultrasound for quantification of arterial structures in human neonates – Comparison with conventional high resolution vascular ultrasound imaging. *Atherosclerosis* 2015;239:523-7.

Touboul PJ, Hennerici MG, Meairs S, Adams H, Amarenco P, Bornstein N, Csiba L, Desvarieux M, Ebrahim S, Hernandez Hernandez R, Jaff M, Kownator S, Naqvi T, Prati P, Rundek T, Sitzer M, Schminke U, Tardif JC, Taylor A, Vicaute E, Woo KS. Mannheim carotid intima-media thickness and plaque consensus (2004-2006-2011). An update on behalf of

the advisory board of the 3rd, 4th and 5th watching the risk symposia, at the 13th, 15th and 20th European Stroke Conferences, Mannheim, Germany, 2004, Brussels, Belgium, 2006, and Hamburg, Germany, 2011. *Cerebrovasc Dis* 2012;34:290-6.

Vatanen A, Sarkola T, Ojala TH, Turanlahti M, Jahnukainen T, Saarinen-Pihkala UM, Jahnukainen K. Radiotherapy-related arterial intima thickening and plaque formation in childhood cancer survivors detected with very-high resolution ultrasound during young adulthood. *Pediatric Blood & Cancer* 2015;62:2000-6.

Velican D, Velican C. Comparative study on age-related changes and atherosclerotic involvement of the coronary arteries of male and female subjects up to 40 years of age. *Atherosclerosis* 1981;38:39-50.

Wendelhag I, Gustavsson T, Suurkula M, Berglund G, Wikstrand J. Ultrasound measurement of wall thickness in the carotid artery: fundamental principles and description of a computerized analysing system. *Clin Physiol* 1991;11:565-77.

Wikstrand J. Methodological considerations of ultrasound measurement of carotid artery intima-media thickness and lumen diameter. *Clin Physiol Funct Imaging* 2007;27:341-5.

Wilens SL. The Nature of Diffuse Intimal Thickening of Arteries. *Am J Pathol* 1951;27:825-39.

Table 1.

<b>Histology vs. VHRU</b>	<b>Histology</b>			<b>VHRU</b>		<i>Mean</i>			
	<i>N</i>	<i>Mean</i>	<i>SD</i>	<i>Mean</i>	<i>SD</i>	<i>Difference</i>	<i>LOA 95CI%</i>	<i>CV%</i>	<i>ICC (95%CI)</i>
Intima thickness [mm]	28	0.125	0.045	0.132	0.050	0.007	-0.042;0.057	19.7	0.923 (0.833;0.964)
Intima-media thickness [mm]	37	0.255	0.097	0.243	0.086	-0.012	-0.086;0.064	15.1	0.955 (0.912;0.977)

Table 1. Comparison of histology and VHRU measurements. SD -standard deviation; LOA – 95% limits of agreement; CV% - coefficient of variation (%); ICC – intraclass correlation; 95%CI – 95% confidence interval.

Table 2.

Intra-observer	N	Mean	SD	Mean		LOA 95CI%	CV%	ICC (95% CI)
				Difference				
Intima thickness [mm]	25	0.140	0.047	-0.011		-0.053; 0.032	15.7	0.946 (0.877; 0.976)
Intima-media thickness [mm]	31	0.246	0.090	-0.006		-0.085; 0.071	16.1	0.951 (0.898; 0.976)
<b>Inter-observer</b>								
Intima thickness [mm]	25	0.136	0.044	0.012		-0.040;0.065	19.9	0.872 (0.773; 0.943)
Intima-media thickness [mm]	31	0.240	0.085	0.025		-0.075;0.125	21.2	0.857 (0.704; 0.931)

Table 2. Intra- and inter-observer comparisons. SD – standard deviation; LOA95% - 95% limits of agreement; CV% - Coefficient of variation (%); ICC – Intraclass correlation; 95%CI – 95% confidence interval.

## Figures

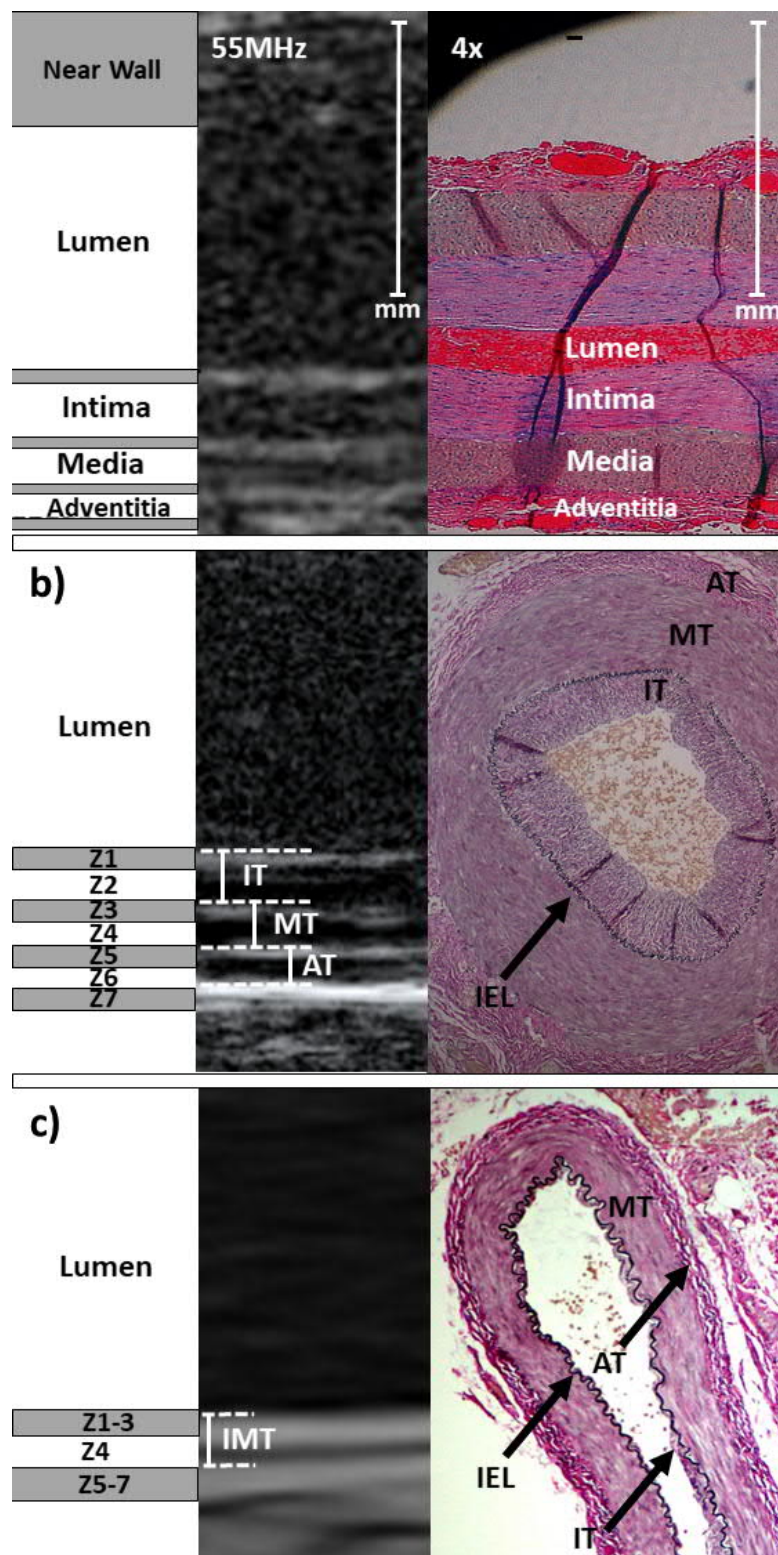


Figure 1. Images of temporal artery histology and corresponding in vivo VHRU (55MHz). a) 4x optic zoom HE-stain histology and VHRU image with identical scales. Note the three layers of the vascular wall is separately distinguishable in the VHRU image. b) Thick

temporal artery wall (VEG-stain) and VHRU image. Note the far wall is separated in to seven different zones displaying a four-line pattern in the VHRU image. Zone 1: echodense lumen intima interface; Zone 2: echolucent intima core; Zone 3: echodense intima-media interface; Zone 4: echolucent media; Zone 5: echodense media-adventitia interface; Zone 6: echolucent adventitia; Zone 7: echodense adventitia-vascular border. c) Thin intima of temporal artery wall on histology represented by the inner surface of the stained IEL (VEG-stain) and on the corresponding VHRU image showing fused echo zones 1-3. echolucent media zone 4. and echodense media vascular wall interface (zones 5-7). Note that the adventitial layer thickness in this temporal artery lacks sufficient thickness to be measured. AT – Adventitia; IT – Intima; MT – Media; IMT – Intima-media thickness; IEL – Internal elastic lamina.



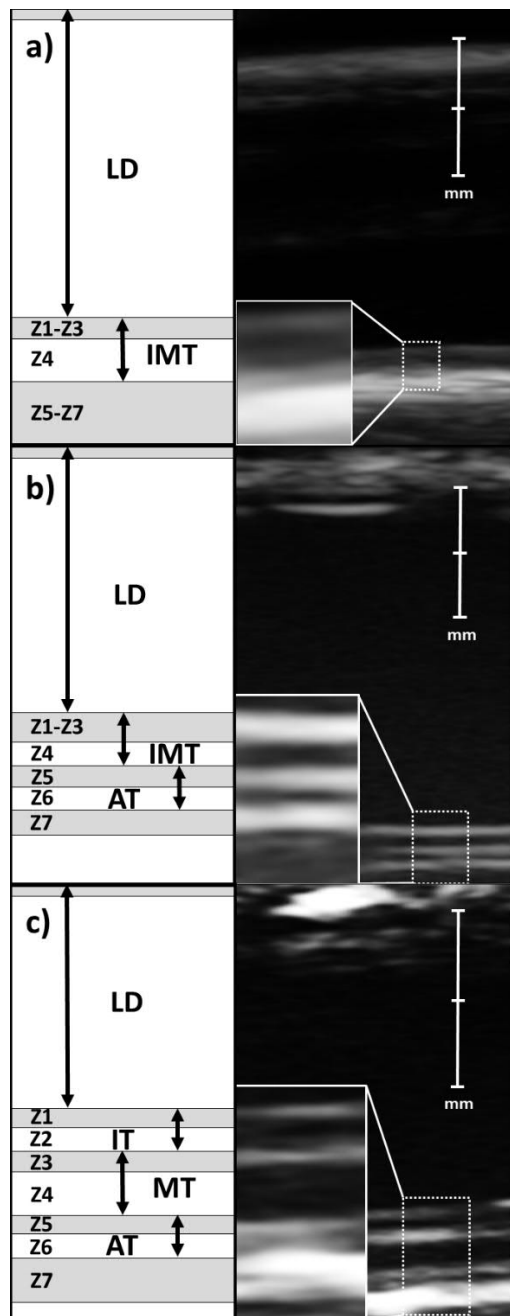


Figure 2. Schematic and VHRU images showing a) the double-line pattern seen in the elastic common carotid artery (52yrs. Female, 25MHz) with fused echo zones 1-3 and 5-7 b) triple-line pattern seen in a muscular radial artery (53yrs. Female, 55MHz) with separation of the media-adventitial echo zones into three separate echo zones (5-7). and c) the distinct four-line pattern in an aged muscular radial artery (male. 77yrs, 55MHz). with separation of the intima-media echo zones (1-3) and media-adventitia echo zones (5-7). Please note the interfaces applied in the leading-to-leading edge measurement technique for far wall IT, IMT, and AT.

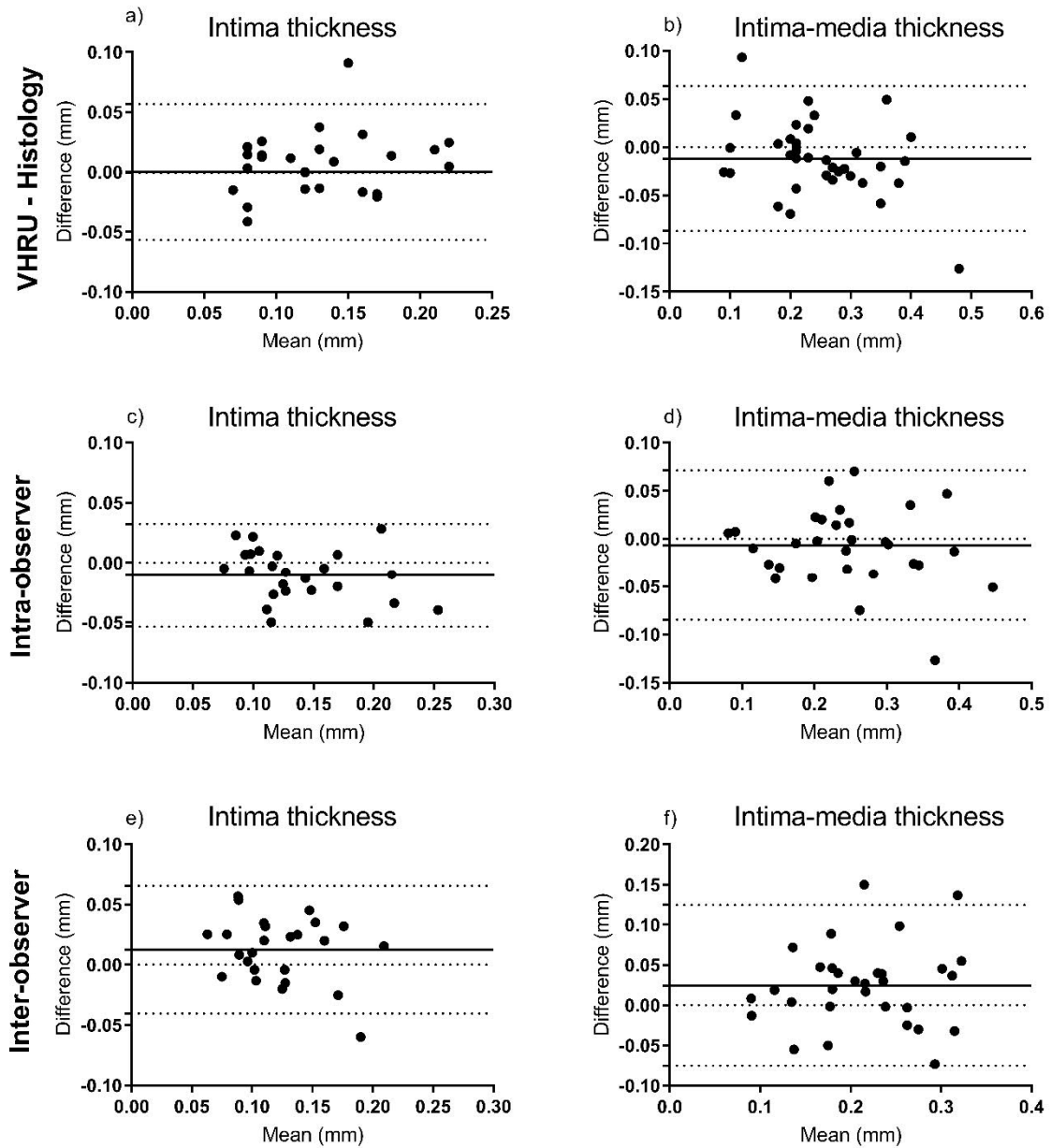
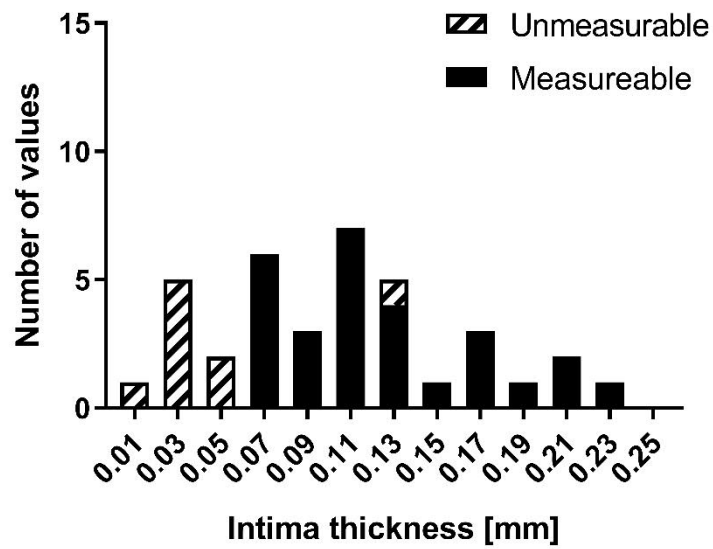
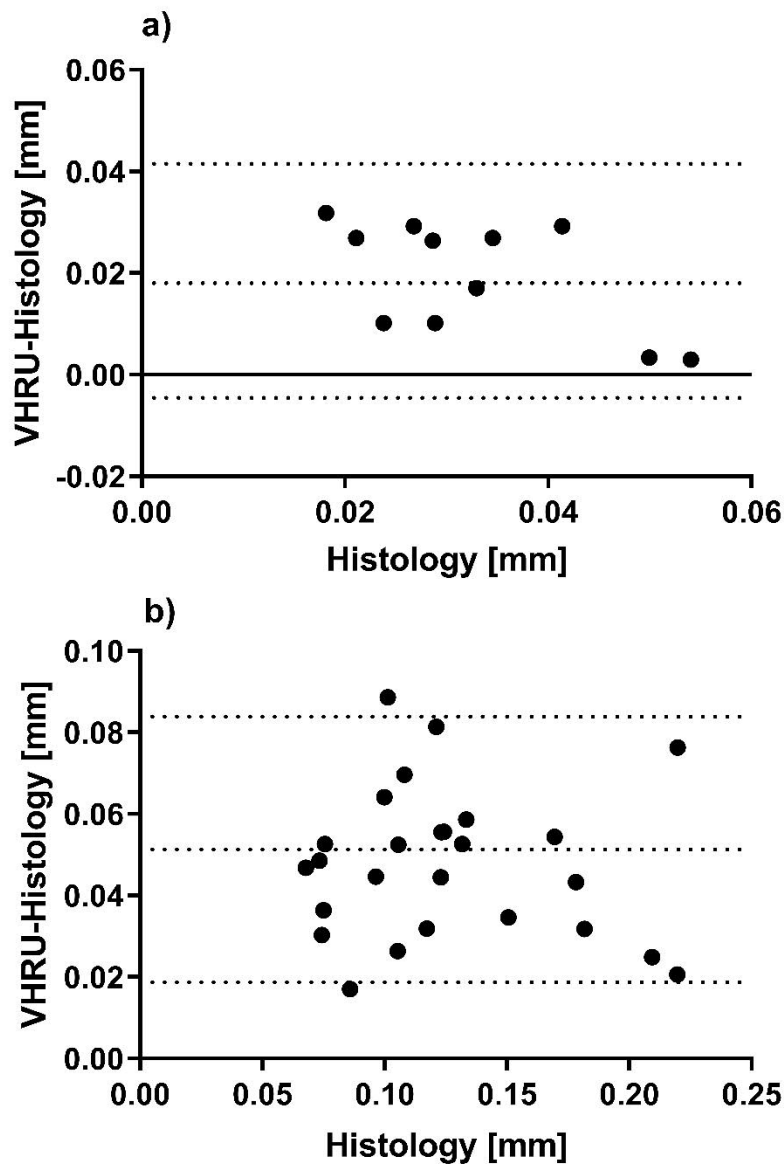


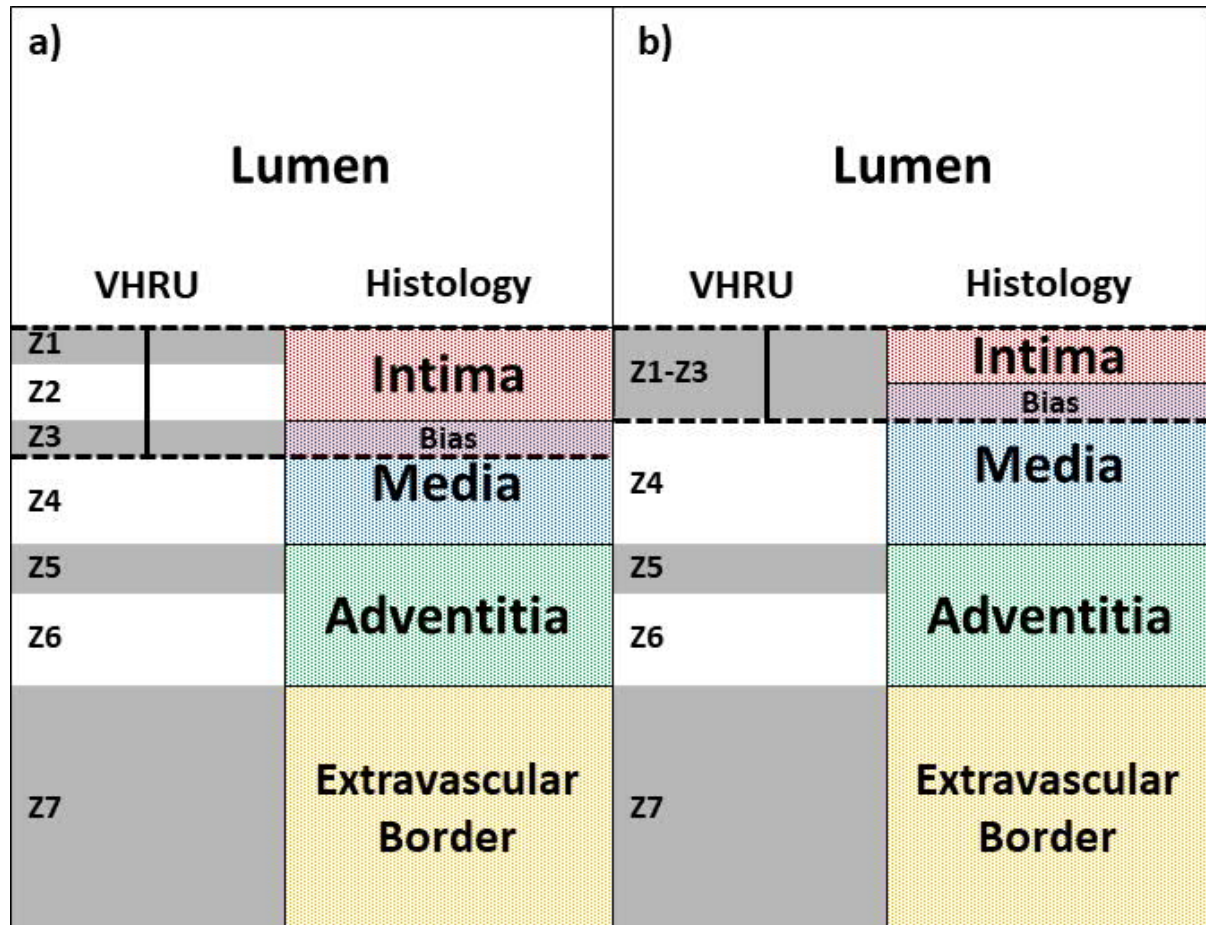
Figure 3. Bland Altman plots of very-high resolution ultrasound (55MHz VHRU) vs. Histology accuracy a) and b). intra-observer agreements c) and d) and inter-observer agreements e) and f) for temporal artery intima (IT) and intima-media thickness (IMT).

**Supplemental Figure Legends**

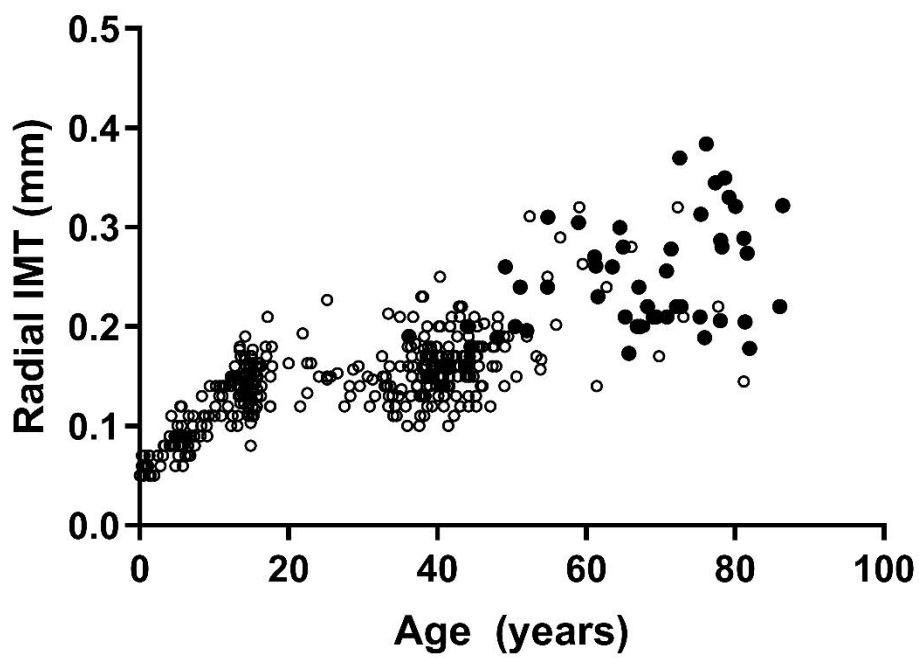
Supplemental figure 1. Distribution of temporal artery intima thickness (IT) on histology separated into groups according to four-line pattern visibility and measurable IT with 55MHz VHRU.



Supplemental figure 2. Bland-Altman plots comparing very-high resolution ultrasound (VHRU) intima layer thickness (IT) with histology IT using the leading-to-trailing edge measurement technique in the assessment of the VHRU image. a) Leading-to-trailing edge VHRU measurement of blood-intima interface (fused zones 1-3) in arteries with histological IT less than 0.06 mm. b) Leading-to-trailing edge VHRU measurement of the visible IT (zones 1-3 separated in image, measurement from leading-edge of zone 1 to trailing edge of zone 3) in arteries with histological IT 0.06 mm or more. Note the systematic bias in leading-to-trailing edge VHRU IT measurement in comparison to histological IT in both settings.



Supplemental figure 3. Schematic image of how measures using the leading-to-trailing edge method induces bias in a) vessels with a thickened intima (separated zones 1-3). and b) vessels with a thin intima (fused zones 1-3). Note how the bias is equal to Z3 in case a). whereas the bias is influenced by the histological intima artery layer thickness (IT) in b) as Z1-Z3 remains constant and non-related with histological IT variance.



Supplemental figure 4. Radial artery intima-media thickness, from 55 MHz or UHF70 VHRU images, in relation to age including occurrence of visible four-line pattern (black circle) increasing from end of fourth decade of life.

**Supplemental Tables to Sundholm et al Assessing Vascular Intima with Ultrasound**

	Vevo 770	Vevo MD
<i>Release Year</i>	2005	2016
<i>Transducer type</i>	Single Mechanical	Multiple Electrical
<i>Image post-processing</i>	None	Despeckling filter
<i>Multi-focus</i>	No	Yes
<b>Transducers</b>	<b>RMV710B</b>	<b>UHF22</b>
<i>Center transmit</i>	25MHz	15MHz
<i>Frequency range</i>	12-38MHz	10-22MHz
<i>Axial Resolution</i>	35.7µm/px	44.8µm/px
<i>Penetration</i>	22.5mm	38.4mm
<b>Transducers</b>	<b>RMV712</b>	<b>UHF48</b>
<i>Center transmit</i>	35MHz	30MHz
<i>Frequency range</i>	17-53MHz	20-46MHz
<i>Axial Resolution</i>	19.6µm/px	21.8µm/px
<i>Penetration</i>	13.0mm	23.5mm
<b>Transducers</b>	<b>RMV708</b>	<b>UHF70</b>
<i>Center transmit</i>	55MHz	50MHz
<i>Frequency range</i>	22-83MHz	29-71MHz
<i>Axial Resolution</i>	15.6µm/px	12.3µm/px
<i>Penetration</i>	8.0mm	10.0mm

Supplemental table 1. Comparison of the two ultrasound systems and transducers used in this study.



<b>Sample</b>	<b>1.</b>	<b>2.</b>	<b>3.</b>	<b>4.</b>
<i>N</i>	139	39	24	178
<i>Age (Range)</i>	9.0 (0.0-17.8)	14.6 (13.2-16.3)	29.8 (20.0-46.3)	40.3 (27.5-50.6)
<i>Female (%)</i>	58 (42%)	19 (49%)	0 (0%)	178 (100%)
<i>BMI Z-score</i>				
<i>(Range)</i>	0.0 (-2.1-2.9)	0.57 (-1.3-2.4)		
<i>BMI (Range)</i>	18.3 (11.7-31.9)	22.5 (16.7-34.2)	24.7 (20.4-28.6)	32.2 (17.7-45.1)
<i>Diabetes (%)</i>	0 (0%)	39 (100%)	0 (0%)	10 (5.6%)
<i>Hypertension (%)</i>	0 (0%)	0 (0%)	0 (0%)	19 (10.7%)

Supplemental table 2. Subject characteristics for the four convenience samples included to assess the presence of the four-line pattern in relation to age.

Histology				VHRU					
<i>Histology vs.</i>				<i>Mean</i>					
<i>VHRU</i>	<i>N</i>	<i>Mean</i>	<i>SD</i>	<i>Mean</i>	<i>SD</i>	<i>Difference</i>	<i>LOA 95CI%</i>	<i>CV%</i>	<i>ICC (95%CI)</i>
									0.867
Vevo 770, 55 MHz	15	0.116	0.041	0.127	0.049	0.011	-0.049;0.069	24,9	(0.603;0.955)
									0.971
Vevo MD, 50 MHz	12	0.135	0.050	0.139	0.053	0.004	-0.030;0.037	12,6	(0.901;0.992)

Supplemental table 3. Comparison of histology and VHRU measurements using the 55 MHz Vevo 770 and 50 MHz Vevo MD, respectively. SD -standard deviation; LOA – 95% limits of agreement; CV% - coefficient of variation (%); ICC – intraclass correlation; 95%CI – 95% confidence interval.

Dependent variable	Adjusted R2	Model p-value
Intima thickness [ $\mu\text{m}$ ]	0.404	<0.001
Independent variables	Beta	p-value
Constant	-36.6	0.464
Age [years]	1.7	0.038
Hypertension [yes=1 no=0]	42.2	0.02
Diabetes [yes=1 no=0]	34.2	0.044
Hypercholesterolemia [yes=1 no=0]	-26.7	0.141
Smoking [10 pack years]	6.8	0.069

Supplemental table 4. Linear regression model assessing effects of age and cardiovascular risk factors on histological intima thickness in the GCA sample (n=37).

<b>Vascular parameters</b>	<b>Mean (SD)/N(%)</b>
<i>Carotid Artery 25MHz [N=30]</i>	
Intima-media thickness	
[mm]	0.65 (0.12)
Plaque presence	12 (40.0%)
Multiple plaques	9 (30.0%)
<i>Radial Artery 55Mhz [N=34]</i>	
Intima-media thickness	
[mm]	0.23 (0.06)
Adventitia thickness [mm]	0.11 (0.03)
4-line pattern	23 (67.6%)
Intima thickness	0.11 (0.06) <sup>a</sup>
<i>Brachial Artery 35MHz [N=28]</i>	
Lumen Diameter [mm]	3.49 (0.59)
Intima-media thickness	
[mm]	0.30 (0.07)
Adventitia thickness [mm]	0.17 (0.08)
4-line pattern	1(3.5%)

Supplemental table 5. Vascular dimensions. intimal thickening and plaque findings of carotid. brachial and radial arteries in the GCA sample. <sup>a</sup> - calculated from arteries with visible four-line pattern.

Dependent variable			Model p-value
Visible 4-line [0 = no, 1 = yes]			<0.001
Independent variables	Beta (SE)	OR (95%CI)	p-value
Constant	-10.326 (1.137)		
Age [10 years]	1.124 (0.189)	3.077 (2.125-4.456)	<0.001
IMT [0.1mm]	1.153 (0.532)	4.629 (1.631-13.140)	0.004

Supplemental table 6. Logistic regression model of visible radial artery far wall four-line pattern using VHRU. SE – standard error; OR – odds ratio; 95%CI – 95% confidence interval.